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What is claimed is:

- 1.. A transgenic mouse comprising:
 - a) a disrupted H2 class I gene;
 - b) a disrupted H2 class II gene; and
 - c) a functional HLA class I or class II transgene.
- 2. A transgenic mouse comprising:
 - a) a disrupted H2 class I gene;
 - b) a disrupted H2 class II gene;
 - c) a functional HLA class I transgene; and
 - d) a functional HLA class II transgene.
- 3. The transgenic mouse according to claim 2, wherein the HLA class I transgene is an HLA-A2 transgene and the HLA class II transgene is an HLA-DR1 transgene.
- 4. The transgenic mouse according to claim 3, wherein the HLA-A2 transgene comprises the HLA-A2 sequence provided in the sequence listing and the HLA-DR1 transgene comprises the HLA-DR1 sequence provided in the sequence listing.

- 5. A transgenic mouse deficient for both H2 class I and class II molecules, wherein the transgenic mouse comprises a functional HLA class I transgene and a functional HLA class II transgene.
- 6. The transgenic mouse according to claim 5, having the genotype $HLA-A2^{+}HLA-DR1^{+}\beta 2m^{\circ}IA\beta^{\circ}$.
- 7. The transgenic mouse according to claim 6, wherein the HLA-A2 transgene comprises the HLA-A2 sequence provided in the sequence listing and the HLA-DR1 transgene comprises the HLA-DR1 sequence provided in the sequence listing.
- 8. A method of simultaneously identifying the presence of one or more epitopes in a candidate antigen or group of antigens, wherein the epitope elicits a specific humoral response, a TH HLA-DR1 restricted response, and/or a CTRL HLA-A2 restricted response, the method comprising:
 - a) administering the candidate antigen or group of candidate antigens to the mouse of claim 3 or claim 6;
 - b) assaying for a specific humoral response in the mouse to the antigen;
 - c) assaying for a TH HLA-DR1 restricted response in the mouse to the antigen; and

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d) assaying for a CTRL HLA-A2 restricted response in the mouse to the antigen; wherein,

observation of a specific humoral response in the mouse to the antigen identifies an epitope which elicits a humoral response in the antigen;

observation of a TH HLA-DR1 restricted response in the mouse to the antigen identifies an epitope which elicits a TH HLA-DR1 restricted response in the antigen; and

observation of a CTRL HLA-A2 restricted response in the mouse to the antigen identifies an epitope which elicits a CTRL HLA-A2 restricted response in the antigen.

9. The method of claim 8, further comprising assaying for a Th1-specific response in the mouse to the antigen and assaying for a Th2-specific response in the mouse to the antigen; wherein

observation of a Th1-specific response in the mouse to the antigen identifies an epitope which elicits a Th1-specific response in the mouse to the antigen; and

observation of a Th2-specific response in the mouse to the antigen identifies an epitope which elicits a Th2-specific response in the mouse to the antigen.

- 10. A method of identifying the presence of an HLA DR1-restricted T helper epitope in a candidate antigen or group of candidate antigens, the method comprising:
 - a) administering the candidate antigen or group of candidate antigens to the mouse of claim 3 or claim 6; and

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b) assaying for a TH HLA-DR1 restricted T helper epitope response in the mouse to the antigen; wherein,

observation of a TH HLA-DR1 restricted T helper epitope response in the mouse to the antigen identifies an epitope which elicits a TH HLA-DR1 restricted T helper epitope response in the antigen.

- 11. An isolated antigen comprising an HLA DR1-restricted T helper epitope identified by the method of claim 10.
- 12. The isolated antigen of claim 11, wherein the antigen further comprises an epitope which elicits a humoral response and/or an epitope which elicits a CTRL HLA-A2 restricted response.
- 13. The isolated antigen of claim 11, wherein the antigen comprising an HLA DR1-restricted T helper epitope comprises a polypeptide.
- 14. The isolated antigen of claim 11, wherein the antigen comprising an HLA DR1-restricted T helper epitope comprises a polynucleotide.

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- 15. The isolated antigen of claim 14, wherein the antigen comprising an HLA DR1-restricted T helper epitope comprises, DNA, RNA, or DNA and RNA.
- 16. A method of identifying the presence of an HLA-A2-restricted T cytotoxic (CTL) epitope in a candidate antigen or group of candidate antigens, the method comprising:
 - a) administering the candidate antigen or group of candidate antigens to the mouse of claim 3 or claim 6; and
 - b) assaying for an HLA-A2-restricted T cytotoxic (CTL) response in the mouse to the antigen or group of antigens; wherein,

observation of an HLA-A2-restricted T cytotoxic (CTL) response in the mouse to the antigen or group of antigens identifies an epitope which elicits a an HLA-A2-restricted T cytotoxic (CTL) response in the antigen group of antigens.

- 17. An isolated antigen comprising an HLA-A2-restricted T cytotoxic (CTL) epitope identified by the method of claim 16.
- 18. The isolated antigen of claim 17, wherein the antigen further comprises an epitope which elicits a humoral response and/or an epitope which elicits a TH HLA-DR1 restricted T helper epitope response.

- 19. The isolated antigen of claim 17, wherein the antigen comprising an HLA-A2-restricted T cytotoxic (CTL) epitope comprises a polypeptide.
- 20. The isolated antigen of claim 17, wherein the antigen comprising an HLA-A2-restricted T cytotoxic (CTL) epitope comprises a polynucleotide.
- 21. The isolated antigen of claim 20, wherein the antigen comprising an HLA-A2-restricted T cytotoxic (CTL) epitope comprises, DNA, RNA, or DNA and RNA.
- 22. A method of comparing the efficiency of T-helper cell response induced by two or more vaccines, the method comprising:
 - a) administering a first candidate vaccine to a mouse of claim 3 or claim 6 and measuring the T-helper cell response induced in the mouse by the first candidate vaccine;
 - b) administering a second candidate vaccine to a mouse of claim 3 or claim 6 and measuring the T-helper cell response induced in the mouse by the second candidate vaccine;
 - c) administering each additional candidate vaccine to be compared to a mouse of claim 3 or claim 6 and measuring the T-helper cell response induced in the mouse by each additional candidate vaccine to be compared; and

- d) determining the efficiency of each candidate vaccine to induce a

 T-helper cell response by comparing the T-helper cell responses to each of the
 vaccines to be compared with each other.
- 23. The method of claim 22, wherein the T-helper cell response is an HLA-DR1 restricted response.
- 24. A method of comparing the efficiency of T cytotoxic cell response induced by two or more vaccines, the method comprising:
 - a) administering a first candidate vaccine to a mouse of claim 3 or
 claim 6 and measuring the T cytotoxic cell response induced in the mouse by the
 first candidate vaccine;
 - b) administering a second candidate vaccine to a mouse of claim 3 or claim 6 and measuring the T cytotoxic cell response induced in the mouse by the second candidate vaccine;
 - c) administering each additional candidate vaccine to be compared to a mouse of claim 3 or claim 6 and measuring the T cytotoxic cell response induced in the mouse by each additional candidate vaccine to be compared; and
 - d) determining the efficiency of each candidate vaccine to induce a T cytotoxic cell response by comparing the T cytotoxic cell responses to each of the vaccines to be compared with each other.

- 25. The method of claim 24, wherein the T cytotoxic cell response is an HLA-A2 restricted response.
- 26. A method of simultaneously comparing the efficiency of T-helper cell response and T cytotoxic cell response induced by two or more vaccines, the method comprising:
 - a) administering a first candidate vaccine to a mouse of claim 3 or claim 6 and measuring the T-helper cell response and T cytotoxic cell response induced in the mouse by the first candidate vaccine;
 - b) administering a second candidate vaccine to a mouse of claim 3 or claim 6 and measuring the T-helper cell response and T cytotoxic cell response induced in the mouse by the second candidate vaccine;
 - c) administering each additional candidate vaccine to be compared to a mouse of claim 3 or claim 6 and measuring the T-helper cell response and T cytotoxic cell response induced in the mouse by each additional candidate vaccine to be compared; and
 - d) determining the efficiency of each candidate vaccine to induce a Thelper cell response and T cytotoxic cell response by comparing the T-helper cell
 response and T cytotoxic cell response to each of the vaccines to be compared
 with each other.

- 27. The method of claim 26, wherein the T-helper cell response is an HLA-DR1 restricted response, and wherein the T cytotoxic cell response is an HLA-A2 restricted response.
- 28. A method of simultaneously determining the humoral response, the T-helper cell response, and the T cytotoxic cell response of a mouse following its immunization with an antigen or a vaccine comprising one or more antigens, the method comprising:
 - a) administering the antigen or the vaccine comprising one or more antigens to a mouse of claim 3 or claim 6;
 - b) assaying for a specific humoral response in the mouse to the antigen or vaccine comprising one or more antigens;
 - c) assaying for a T-helper cell response in the mouse to the antigen or vaccine comprising one or more antigens; and
 - d) assaying for a T cytotoxic cell response in the mouse to the antigen or vaccine comprising one or more antigens.
- 29. The method of claim 28, wherein the T-helper cell response is a TH HLA-DR1 restricted response.
- 30. The method of claim 28, wherein the T cytotoxic cell response is a CTRL HLA-A2 restricted response.

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31. A method of optimizing two or more candidate vaccine compositions for administration to a human, based on preselected criteria, the method comprising:

simultaneously determining the humoral response, the T-helper cell response, and the T cytotoxic cell response of a mouse following its immunization with the two or more candidate vaccine compositions, according to claim 28; and

selecting an optimized vaccine by applying preselected criteria to the results.

- 32. The method according to claim 31, wherein the two or more candidate vaccines differ only in the ratio of antigen to adjuvant present in the vaccine.
- 33. The method according to claim 31, wherein the two or more candidate vaccines differ only in the type of adjuvant present in the vaccine.
- 34. A method of determining whether a vaccine poses a risk of induction of an autoimmune disease when administered to a human, the method comprising:
 - a) administering the vaccine to a mouse of claim 3 or claim 6; and
- b) assaying for an autoimmune response in the mouse; wherein, observation of an autoimmune response in the mouse indicates that the vaccine poses a risk of induction of an autoimmune disease when administered to a human.
 - 35. An isolated transgenic mouse cell comprising:
 - a) a disrupted H2 class I gene;
 - b) a disrupted H2 class II gene; and

- c) a functional HLA class I or class II transgene.
- 36. An isolated transgenic mouse cell comprising:
 - a) a disrupted H2 class I gene;
 - b) a disrupted H2 class II gene;
 - c) a functional HLA class I transgene; and
 - d) a functional HLA class II transgene.
- 37. The transgenic mouse cell according to claim 36, wherein the HLA class I transgene is an HLA-A2 transgene and the HLA class II transgene is an HLA-DR1 transgene.
- 38. The transgenic mouse cell according to claim 37, wherein the HLA-A2 transgene comprises the HLA-A2 sequence provided in the sequence listing and the HLA-DR1 transgene comprises the HLA-DR1 sequence provided in the sequence listing.
- 39. An isolated transgenic mouse cell deficient for both H2 class I and class II molecules, wherein the transgenic mouse cell comprises a functional HLA class I transgene and a functional HLA class II transgene.
- 40. The transgenic mouse cell according to claim 39, having the genotype HLA-A2⁺HLA-DR1⁺β2m°IAβ°.

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The transgenic mouse cell according to claim 40, wherein the HLA-A2 41. transgene comprises the HLA-A2 sequence provided in the sequence listing and the HLA-DR1 transgene comprises the HLA-DR1 sequence provided in the sequence listing.